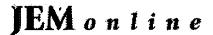


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Clin. Diagn. Lab. Immunol., Nov 1996; 3: 746 - 752. [Abstract] [PDF]



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Diagnosis of contagious caprine pleuropneumonia by detection and identification of Mycoplasma capriculum subsp. capripneumoniae by PCR and restriction enzyme analysis

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F Bonnet, C Saillard, JM Bove, RH Leach, DL Rose, GS Cottew, and JG

DNA relatedness between field isolates of Mycaplasma F38 group, the agent of contagious caprine pleuropneumonia, and strains of Mycopiasma capricolum

Int. J. Syst. Bacteriol., Jul 1993; 43: 597 - 602. [Abstract]

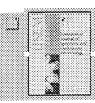


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Mycoplasma from contagious caprine pleuropneumonia Ann. N.Y. Acad. Sci., Jul 1967; 143; 298 - 304.



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Relationships between members of the Mycoplasma mycoides cluster as shown by DNA probes and sequence analysis Int. J. Syst. Bacteriol., Oct 1992; 42: 593 - 601. [Abstract]



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Vaccination against Contagious Bovine Pleuropneumonia and the Use of Molecular Tools in Epidemiology Ann. N.Y. Acad. Sci., Jun 1998; 849: 146 - 151.

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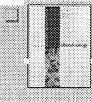
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Bertil Pettersson, Göran Bölske, François Thiaucourt, Mathlas Uhlén, and Karl-Erik Johansson

Molecular Evolution of *Mycoplasma capricolum* subsp. capripneumoniae Strains, Based on Polymorphisms in the 16S rRNA Genes



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X Cheng, J Nicolet, R Miserez, P Kuhnert, M Krampe, T Pilloud, EM Abdo, C Griot, and J Frey

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A Mycoplasma strain F38 growth-inhibiting monoclonal antibody

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Infect. Immun., Apr 1995; 63: 1415 - 1420. [Abstract] [PDF]



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CR Bascunana, JG Mattsson, G Bolske, and KE Johansson

Characterization of the 16S rRNA genes from Mycopiasma sp. strain

F38 and development of an identification system based on PCR

J. Bacteriol., May 1994; 176: 2577 - 2586. [Abstract]



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Proposal for designation of F38-type caprine mycoplasmas as Mycoplasma capricolum subsp. capripneumoniae subsp. nov. and consequent obligatory relegation of strains currently classified as M. capricolum (Tully, Barile, Edward, Theodore, and Erno 1974) to an additional new subspecies, M. capricolum subsp. capricolum subsp. nov

Int. J. Syst. Bacteriol., Jul 1993; 43: 603 - 605. [Abstract]



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Acute respiratory, circulatory and pathological changes in the calf after intravenous injections of the galactan from Mycoplasma mycoides subsp. mycoides

J. Med. Microbiol., Nov 1976; 9: 379 - 391. [Abstract]



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WA, Clyde, Jr. and FW Denny

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Pediatrics, Oct 1967; 40: 669 - 684. [Abstract]



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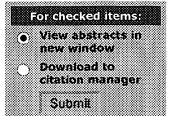
SJ Sussman, RL Magoffin, EH Lennette, and J Schieble

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Pediatrics, Oct 1966; 38: 571 - 577. [Abstract]

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WC Lai, SP Pakes, K Ren, YS Lu, and M Bennett

Therapeutic effect of DNA immunization of genetically susceptible mice infected with virulent Mycoplasma pulmonis

J Immunol 1997 158: 2513-2516. [Abstract] [PDF]

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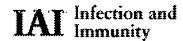
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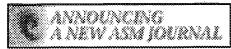
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Immunoblot analyses of chimpanzee sera after infection and after immunization and challenge with Mycoplasma pneumoniae

G Franzoso, PC Hu, GA Meloni and MF Barile

Laboratory of Mycoplasma, Food and Drug Administration, Bethesda, Maryland 20892.

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or 4-month study from 16 chimpanzees were examined by immunoblot analyses to identify the immunogenic components of Mycoplasma pneumoniae. Six experimentally infected chimpanzees showed significant signs of overt disease, including cough, pharyngitis, rhinitis, fever, and loss of appetite. The sera of these infected chimpanzees recognized from 17 to 20 protein bands. Two control chimpanzees that were not inoculated were included in the study. Three chimpanzees immunized with a formalininactivated OSU-1A vaccine and three chimpanzees immunized with an experimental acellular vaccine showed minimal signs of disease on challenge. After challenge, the serum immunoblot responses of the immunized chimpanzees were similar to those of the infected chimpanzees. Before challenge, the sera of two previously infected chimpanzees recognized protein bands of 169 (which comigrated with the P1 adhesin), 148, 130, 117, 86, 61, 44, 35, 30, and 29 kDa. After challenge, the previously infected chimpanzees showed the most intense serum immunoblot responses and were most protected against colonization and disease. The sera from each of the 16 chimpanzees examined recognized a large number of immunogenic components, and the serum immunoblot responses were virtually identical to those of patients. Sera from each chimpanzee and patient recognized 169-, 148-, 130-, 117-, 86-, 44-, and 35-kDa bands and many of them recognized 67-, 63-, 61-, 56-, 32-, 30-, and 29-kDa protein bands.

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- SVENSTRUP, H. F., NIELSEN, P. K., DRASBEK, M., BIRKELUND, S., CHRISTIANSEN, G. (2002). Adhesion and inhibition assay of Mycoplasma genitalium and M. pneumoniae by immunofluorescence microscopy. *J Med Microbiol* 51: 361-373 [Abstract] [Full Text]
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TI Xerovac: an ultra rapid method for the dehydration and **preservation** of live attenuated Rinderpest and Peste des Petits ruminants **vaccines**.

AU Worrall E E; Litamoi J K; Seck B M; Ayelet G

SO VACCINE, (2000 Nov 22) 19 (7-8) 834-9. Journal code: 8406899. ISSN: 0264-410X.

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TI [A device for safe handling of **vacuum-dried** microbes (author's transl)].

Eine Einrichtung zum gefahrlosen Umgang mit vacuumgetrockneten Mikroben.

AU Eyer H; Schmidt M

- SO. ZENTRALBLATT FUR BAKTERIOLOGIE, PARASITENKUNDE, INFEKTIONSKRANKHEITEN UND HYGIENE. ERSTE ABTEILUNG ORIGINALE. REIHE A: MEDIZINISCHE MIKROBIOLOGIE UND PARASITOLOGIE, (1975) 230 (4) 534-7.

 Journal code: 0331570. ISSN: 0300-9688.
- L11 ANSWER 3 OF 12 CAPLUS COPYRIGHT 2003 ACS
- TI Scalable long-term shelf **preservation** of sensitive **biological** solutions and suspensions

IN Bronshtein, Victor

- SO U.S., 13 pp., Cont.-in-part of U.S. Ser. No. 785,472, abandoned. CODEN: USXXAM
- L11 ANSWER 4 OF 12 CAPLUS COPYRIGHT 2003 ACS
- TI Preservation and formulation of bioactive materials for storage and delivery in hydrophobic carriers
- IN Bronshtein, Victor
- SO PCT Int. Appl., 36 pp. CODEN: PIXXD2
- L11 ANSWER 5 OF 12 CAPLUS COPYRIGHT 2003 ACS
- TI Method for the preservation of viruses and mycoplasma
- IN Worrall, Eric Edward
- SO PCT Int. Appl., 24 pp. CODEN: PIXXD2
- L11 ANSWER 6 OF 12 CAPLUS COPYRIGHT 2003 ACS
- TI Method and composition for preserving viruses
- IN Kovesdi, Imre; Ransom, Stephen C.
- SO PCT Int. Appl., 19 pp. CODEN: PIXXD2
- L11 ANSWER 7 OF 12 CAPLUS COPYRIGHT 2003 ACS
- TI **Preservation** of sensitive **biological** samples by vitrification
- IN Bronshtein, Victor
- SO PCT Int. Appl., 26 pp. CODEN: PIXXD2
- L11 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2003 ACS
- TI Protection of proteins and the like
- IN Roser, Bruce Joseph
- SO PCT Int. Appl., 41 pp. CODEN: PIXXD2
- L11 ANSWER 9 OF 12 PROMT COPYRIGHT 2003 Gale Group

- TI A spoonful of sugar.
- AU Haydon, Colette
- SO Soap Perfumery & Cosmetics, (June 2000) Vol. 73, No. 6, pp. 39. ISSN: 0037-749X.
- L11 ANSWER 10 OF 12 PROMT COPYRIGHT 2003 Gale Group
- TI Universal Preservation Technologies Awarded Office of Naval Research Contract for the Preservation of Certain Mammalian Cells.
- SO Business Wire, (10 Apr 2000) pp. 1339.
- L11 ANSWER 11 OF 12 PROMT COPYRIGHT 2003 Gale Group
- TI Trhalose Boosts Prospects for Improved Biopharmaceuticals and Vaccines
 New applications for QT-4 formulation are found in drug & vaccine delivery & in food preservation
- SO Genetic Engineering News, (15 Mar 1995) pp. 10. ISSN: 1270-6377.
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- TI A NEW TECHNIQUE OFFERING ROOM TEMPERATURE STABLE BIOLOGICS
- SO Pharmaceutical Manufacturing Review, (Dec 1994) pp. 34. ISSN: 0955-3894.



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    of industrial scale biol. solns. and suspensions contg.
    biol. active mols., cells and small multicellular
     specimens at ambient temps. by dehydration in amorphous very viscous liq.
     or glass state. The scale up method comprises the primary
     drying step of boiling under vacuum to form a
     mech.-stable foam and a secondary drying step to increase the
     stability. Vitrification can subsequently be achieved by cooling the
     dried material to the storage temp. which is lower than the glass
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            GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KR,
            KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
            NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR,
            TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD,
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RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
PRAI US 1999-166783P
                     P
                           19991122
     This invention relates to formulations comprising biol. samples
AΒ
    preserved as dry glassy powders and hydrophobic
     carriers, the formulations being adapted for the long-term storage and
     delivery of the bioactive materials, in particular viral and
    bacterial vaccines, vectors and cells, at ambient or
    higher temps., and to methods for prepg. these formulations.
     freeze-dried samples of amphotericin B were rehydrated with 40%
     sucrose/vial. Then. The solns. were transferred sterilized glass
    vials for future preservation by drying. B. Before drying, the
    vials were covered with gray Bu slotted rubber stoppers. The vials were
    dried inside a vacuum chamber. Before the vacuum was
     applied the shelf temp. was decreased to 5.degree.. The hydrostatic
    pressure inside the chamber was decreased to 0.5 Torr. The suspension was
    boiled for 30 min. The results of the assay suggested that the loss of
    potency was only detected in those samples dried at the lower temp.
     (25.degree.) and subsequently stored at 40.degree..
    ANSWER 6 OF 12 CAPLUS COPYRIGHT 2003 ACS
L11
ΑN
     2000:401969 CAPLUS
DN
     133:28260
    Method and composition for preserving viruses
TI
IN
     Kovesdi, Imre; Ransom, Stephen C.
    Genvec, Inc., USA
PA
SO
     PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
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     Patent
LΑ
     English
FAN.CNT 1
     PATENT NO.
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            AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
            CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
            IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
            MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
            SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM,
            AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
            DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
            CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
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                                        US 1998-208666
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                           20011004
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        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
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                      A1
                           20020214
                                                            20010531
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                      B2
                           20030204
PRAI US 1998-208666
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AB The present invention provides a method and a compn. for preserving a virus. The virus is placed in a lig. carrier with a stabilizing agent selected from the group consisting of polysorbate 80, L-arginine, polyvinylpyrrolidone, trehalose, and combinations thereof. The liq. compn. can be maintained at a temp.

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above 0 .degree.C for a significant period of time while maintaining a satisfactory degree of viral activity. ANSWER 7 OF 12 CAPLUS COPYRIGHT 2003 ACS L111999:359630 CAPLUS 131:2531 Preservation of sensitive biological samples by vitrification Bronshtein, Victor Universal Preservation Technologies, Inc., USA PCT Int. Appl., 26 pp. CODEN: PIXXD2 Patent English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ---------WO 1997-US21703 19971126 WO 9927071 A1 19990603 W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG AA 19990603 CA 1997-2312233 19971126 CA 2312233 A1 19990615 A1 20000906 AU 1998-54596 19971126 AU 9854596 AU 1990 3.335 EP 1997-948550 EP 1032647 19971126 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI PRAI WO 1997-US21703 19971126 Α This invention discloses methods for the long-term preservation of industrial scale biol. solns. and suspensions contg. biol. active mols., cells and small multicellular specimens at ambient temps. by dehydration in amorphous very viscous liq. or glass state. The scale up method comprises the primary drying step of boiling under vacuum to form a mech.-stable foam and a secondary drying step to increase the stability. Vitrification can subsequently be achieved by cooling the dried material to the storage temp. which is lower than the glass transition temp. THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 6 ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 8 OF 12 CAPLUS COPYRIGHT 2003 ACS 1987:420354 CAPLUS 107:20354 Protection of proteins and the like Roser, Bruce Joseph Quadrant Bioresources Ltd., UK PCT Int. Appl., 41 pp. CODEN: PIXXD2 Patent English FAN.CNT 1

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	EΡ	229810	A1	19870729	EP	1986-904281	19860709
	EΡ	229810	B1	19911016			
		R: AT, BE,	CH, DE	, FR, GB,	IT, LI, I	LU, NL, SE	
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PRAI	GB	1985-17352		19850709			
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AB Sensitive proteins and other macromols., such as enzymes, antibodies, antigens, serum complement, fluorescent proteins, vaccine components, polysaccharides such as agarose, etc., can be preserved by drying at ambient temp. and atm. pressure in the presence of trehalose. A porous matrix impregnated with trehalose is provided as a receiver for a blood or other liq. sample to be dried, e.g. prior to anal. Alk. phosphatase from calf intestine in phosphate-buffered saline was incubated in the wells of an immunoplate overnight. The wells were washed and dried at 37.degree. in the presence or absence of 5% trehalose in distd. water. The enzyme retained full activity on drying in the presence of trehalose, but lost >90% of its activity when dried in the absence of trehalose.

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- TI Physical, chemical and immunological stability of CHO-derived hepatitis B surface antigen (HBsAg) particles.
- AU Diminsky D; Moav N; Gorecki M; Barenholz Y
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- SO PHARMACEUTICAL RESEARCH, (1998 Jul) 15 (7) 1111-6. Journal code: 8406521. ISSN: 0724-8741.
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- AU Anthony P; Ausseil J; Bechler B; Benguria A; Blackhall N; Briarty L G; Cogoli A; Davey M R; Garesse R; Hager R; Loddenkemper R; Marchant R; Marco R; Marthy H J; Perry M; Power J B; Schiller P; Ugalde C; Volkmann D; Wardrop J
- SO JOURNAL OF BIOTECHNOLOGY, (1996 Jun 27) 47 (2-3) 377-93. Journal code: 8411927. ISSN: 0168-1656.
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- AU Grose C; Friedrichs W E; Smith K O
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- SO U.S., 25 pp., Cont.-in-part of U.S. 5,932,223. CODEN: USXXAM
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- TI Formulation of preservation mixtures containing sensitive biologicals to be stabilized for ambient temperature storage by drying
- IN Bronshtein, Victor; Linkowski, Lynn
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- TI Method for preserving biosensor chip
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- SO Jpn. Kokai Tokkyo Koho, 8 pp. CODEN: JKXXAF
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- TI Apparatus for concentrating and **preserving** food products and **biological** cultures, sera, etc.
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- SO Cosmetics & Toiletries Manufacturers & Suppliers, (Dec 1994) pp. 41. ISSN: 0952-519X.
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L12 ANSWER 3 OF 41 MEDLINE MEDLINE AN97141232 DN 97141232 PubMed ID: 8987576 ΤI Preservation of viable biological samples for experiments in space laboratories. Anthony P; Ausseil J; Bechler B; Benguria A; Blackhall N; Briarty L G; ΑU Cogoli A; Davey M R; Garesse R; Hager R; Loddenkemper R; Marchant R; Marco R; Marthy H J; Perry M; Power J B; Schiller P; Ugalde C; Volkmann D; Wardrop J Life Science Department, University of Nottingham, UK. CS JOURNAL OF BIOTECHNOLOGY, (1996 Jun 27) 47 (2-3) 377-93. SO Journal code: 8411927. ISSN: 0168-1656. CY Netherlands DT Journal; Article; (JOURNAL ARTICLE) LΑ FS Biotechnology; Space Life Sciences FΜ 199702 Entered STN: 19970305 ED Last Updated on STN: 19970305 Entered Medline: 19970219 Standard viable preservation methods for biological AΒ samples using low temperatures have been investigated concerning their storage capabilities under higher temperature levels than usual. For a representative set of organism classes (plants, mammalian cells, arthropods and aquatic invertebrates), the minimum appropriate storage conditions have been identified by screening storage temperatures at -196 degrees, -80 degrees, -20 degrees, +4 degrees, +20 degrees/25 degrees C for periods from 2 days to 4 weeks. For storage below 0 degree C, as a typical cryopreservative, dimethylsulfoxide (DMSO) was used. For some samples, the addition of trehalose (as cryopreservative) and the use of a nitrogen atmosphere were investigated. After storage, the material was tested for vitality. The findings demonstrated that acceptable preservation can be achieved under higher storage temperatures than are typically applied. Small, dense cultured plant cells survive for 21 d when moderately cooled (+4 degrees to -20 degrees C); addition of trehalose enhances viability at -20 degrees C. For mammalian cells, the results show that human lymphocytes can be preserved for 3 d at 25 degrees C, 7 d at 4 degrees C and 28 d at -80 degrees C. Friend leukaemia virus transformed cells can be stored for 3 d at 25 degrees C, 14 d at 4 degrees C and 28 d at -80 degrees C. Hybridoma cells can be kept 7 d at 4 degrees C and 28 d at -20 degrees C or -80 degrees C. Model arthropod systems are well preserved for 2 weeks if maintained at lower temperatures that vary depending on the species and/or stage of development; e.g., 12 degrees C for Drosophila imagoes and 4-6 degrees C for Artemia nauplii. For aquatic invertebrates such as sea urchins, embryonic and larval stages can be preserved for several weeks at +6 degrees C, whereas sperm and eggs can best be stored at + 4 degrees C for up to 5 d at maximum. These results enhance the range of feasible space experiments with biological systems. Moreover, for typical terrestrial preservation methods, considerable modification potential is identified. L12 ANSWER 4 OF 41 MEDLINE AN82030023 MEDLINE DN PubMed ID: 6270035 82030023 Cryopreservation of varicella-zoster virions without loss of structural TIintegrity or infectivity. ΑU Grose C; Friedrichs W E; Smith K O

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AI 14604 (NIAID)

INTERVIROLOGY, (1981) 15 (3) 154-60.

Journal code: 0364265. ISSN: 0300-5526. Switzerland · CY Journal; Article; (JOURNAL ARTICLE) DT English LΑ FS Priority Journals EM198112 ED Entered STN: 19900316 Last Updated on STN: 19970203 Entered Medline: 19811215 Varicella-zosterer virions present in infected cells or in a cell-free AΒ state were freeze-dried without loss of structural integrity of infectivity. Generally, yields of greater than 5 log10 foci/ml (infected cells) or greater than 4 log10 PFU/ml (cell-free virus) were recovered from varicella-zoster virus-infected human melanoma cells both before and after lyophilization in phosphate-buffered media containing 0.1-1.0 M sucrose. Virus frozen in solutions lacking sugar had little or no residual infectivity after vacuum sublimation was completed. Visualization by electron microscopy demonstrated large numbers of enveloped virions in the virus preparations lyophilized in media containing sucrose; in marked contrast, virus subjected to freeze-drying in buffered solutions without sugar consisted mainly of naked nucleocapsids. Water analyses by Karl Fischer titration suggested that residual moisture retained by sugar prevented disenvelopment of the varicella-zost virion. ANSWER 6 OF 41 MEDLINE L12MEDLINE AN69250078 DN 69250078 PubMed ID: 5797938 Stabilities of dried suspensions of influenza virus TIsealed in a vacuum or under different gases. Greiff D; Rightsel W A AU APPLIED MICROBIOLOGY, (1969 Jun) 17 (6) 830-5. SO Journal code: 7605802. ISSN: 0003-6919. CY United States DTJournal; Article; (JOURNAL ARTICLE) LΑ English Priority Journals FS EM196909 ED Entered STN: 19900101 Last Updated on STN: 19900101 Entered Medline: 19690917 ANSWER 9 OF 41 CAPLUS COPYRIGHT 2003 ACS L12AN2002:444388 CAPLUS DN 137:10950 Rotavirus vaccine formulations containing a sugar TΙ Burke, Carl J.; Volkin, David B. INPA Merck & Co., Inc., USA U.S., 25 pp., Cont.-in-part of U.S. 5,932,223. SO CODEN: USXXAM DTPatent English LA FAN.CNT 3

APPLICATION NO. DATE KIND DATE PATENT NO. _____ -----US 1999-366616 19990803 B1 20020611 US 6403098 ZA 1997-8586 19970925 ZA 9708586 Α 19980615 US 1997-938260 19970926 19990803 US 5932223 Α WO 2001008495 A1 20010208 WO 2000-US21264 20000803 W: 'AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,

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HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU,
            LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD,
             SE, SG, SI
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                          20020522
                                         EP 2000-955357 20000803
     EP 1206189
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            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL
                     T2 20030212
                                           JP 2001-513242
                                                            20000803
     JP 2003505482
PRAI US 1996-26754P
                       P
                           19960926
    US 1997-46760P
                       Ρ
                           19970516
     US 1997-938260
                      A2
                           19970926
     US 1999-366616
                      Α
                           19990803
     WO 2000-US21264
                      W
                           20000803
     The present invention provides liq. and lyophilized formulations of
AB
     vaccines against rotavirus infection and methods of their prepn.
     The formulations include buffering agents appropriate for oral
     administration of rotavirus vaccines. The formulations also
     include compds. to stabilize the vaccine compns. against loss of
     potency. For example, 1-yr probe stability data were obtained for several
     optimized lyophilized and liq. formulations of G1 and P1 rotavirus at
     various temps. and compared to the stability data of an unoptimized
     formulation, Williams' E (WE) medium/5% sucrose. Optimized liq.
     formulations contq. rotavirus reassortants in WE medium contg. sucrose,
     sodium phosphate, and sodium succinate or sodium citrate showed a
     substantial improvement in stability. Further improvements in storage
     stability were obsd. for lyophilized formulations. With the appropriate
     formulation, the thermostability of rotavirus exceeds that of existing
     live-virus liq. (i.e., OPV) and lyophilized (e.g., measles)
     vaccines. The stabilizing effect of either the
     succinate/phosphate or the citrate/ phosphate buffers offers the potential
     of combining stability enhancement with a gastric neutralization. Liq.
     formulations as well as lyophilized formulations that can be reconstituted
     using this buffer can allow the formulation to be delivered in a single
     administration.
              THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 38
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 10 OF 41 CAPLUS COPYRIGHT 2003 ACS
L12
     2001:396599 CAPLUS
AN
DN
     135:2554
     Formulation of preservation mixtures containing sensitive biologicals to
TI
     be stabilized for ambient temperature storage by drying
IN
     Bronshtein, Victor; Linkowski, Lynn
PA
     Universal Preservation Technologies, Inc., USA
SO
     PCT Int. Appl., 34 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LΑ
FAN.CNT 1
                      KIND DATE
                                          APPLICATION NO.
                                                           DATE
     PATENT NO.
                     ____
                            _____
                      A2
                            20010531
                                          WO 2000-US32261 20001122
ΡI
     WO 2001037656
                      A3
     WO 2001037656
                            20020110
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             CR, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EE, EE, ES, FI, FI, GB,
             GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KR,
             KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
             NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR,
             TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU,
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TJ, TM
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             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                          EP 2000-980766 20001122
                       A2
                           20020821
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
PRAI US 1999-166928P
                           19991122
                      Р
                       W
                            20001122
     WO 2000-US32261
     This invention relates to formulations and methods for preserving
AΒ
     sensitive biologicals, viruses, bacteria and
     eukaryotic cells by drying. More particularly, the invention relates to
     preservation mixts. comprising viruses or cells and
     protectants, including methylated monosaccharides, wherein the mixts. are
     adapted to stabilize these samples during dehydration and subsequent
     storage at ambient and higher temps.
     ANSWER 12 OF 41 CAPLUS COPYRIGHT 2003 ACS
L12
     1997:394163 CAPLUS
AN
     127:23753
DN
     Stabilization of biological materials by drying without freezing
ΤI
     Mattern, Markus; Winter, Gerhard
IN
     Boehringer Mannheim Gmbh, Germany
PA
SO
     Ger. Offen., 32 pp.
     CODEN: GWXXBX
DT
     Patent
LΑ
     German
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                           APPLICATION NO.
                                                            DATE
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                                            ______
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                            19970430
                                           DE 1995-19539574 19951025
ΡI
     DE 19539574
                       A1
                       AA
                            19970501
                                           CA 1996-2235243 19961024
     CA 2235243
                                           WO 1996-EP4627
                       A2
                            19970501
                                                             19961024
     WO 9715288
     WO 9715288
                       A3
                            19970529
             AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK,
             EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR,
             LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM
         RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,
             IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM
     AU 9672984
                            19970515
                                           AU 1996-72984
                                                             19961024
                       A1
                                           ZA 1996-8930
    ZA 9608930
                            19980424
                                                             19961024
                       Α
                                           EP 1996-934811
     EP 857060
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                            19980812
                                                             19961024
     EP 857060
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             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE,
             SI, FI
     CN 1205628
                       Α
                            19990120
                                            CN 1996-199329
                                                             19961024
     BR 9611265
                       Α
                            19990504
                                            BR 1996-11265
                                                             19961024
     JP 11513700
                       T2
                            19991124
                                           JP 1996-516286
                                                             19961024
                       A1
                            20011031
                                           IL 1996-124204
                                                             19961024
     IL 124204
                                           AT 1996-934811
                                                             19961024
     AT 212541
                       Ε
                            20020215
                       T3
                                           ES 1996-934811
     ES 2170274
                            20020801
                                                             19961024
                       C2
                                           RU 1998-109886
                                                             19961024
     RU 2191003
                            20021020
                       Α
                            19980625
                                           NO 1998-1868
                                                             19980424
     NO 9801868
     US 2001055617
                       A1
                            20011227
                                           US 1998-51918
                                                             19980427
PRAI DE 1995-19539574 A
                            19951025
     WO 1996-EP4627
                       W
                            19961024
     A biol., esp. therapeutic, material is stabilized and
AΒ
     preserved by prepg. a soln. of (1) the material, (2) a
     carbohydrate or a zwitterionic compd. with polar residues, and (3) a
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zwitterionic compd. with nonpolar residues, and drying the soln. at a temp. above its f.p. The process does not involve use of elevated temps., can be carried out in conventional lyophilization app., is energy efficient, and is more rapid than freeze drying. Thus, a soln. contg. maltose 50, L-phenylalanine 10, L-arginine 10, polysorbate 80 0.1, and recombinant human G-CSF 0.35 mg/mL (pH 7.4) was sterilized by filtration and 1-mL portions were dispensed into 2-mL vials fitted with lyophilization stoppers and dried isothermally at 20.degree. and reduced pressure for 48 h. The product had a residual water content of 1.16% and a glass transition temp. of 75.degree. The content of native (monomeric) G-CSF was still 99.83% after 13 wk storage at 50.degree.

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L12 ANSWER 13 OF 41 CAPLUS COPYRIGHT 2003 ACS
    1996:685381 CAPLUS
AΝ
DN
    125:309076
    Pharmaceutical composition for preserving recombinant
ΤI
    virus vectors for gene therapy
    Kuma, Hidekazu; Iijima, Osamu; Suzuki, Yousuke
IN
PA
    Hisamitsu Pharmaceutical Co., Inc., Japan
SO
     PCT Int. Appl., 17 pp.
    CODEN: PIXXD2
DT
    Patent
LΑ
    Japanese
FAN.CNT 1
                KIND DATE
                                      APPLICATION NO. DATE
    PATENT NO.
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                         _____
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    WO 9629096 A1 19960926
PΙ
                                        WO 1996-JP652 19960315
        W: AU, CA, CN, JP, KR, US RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE
                                   AU 1996-49544 19960315
                   A1
    AU 9649544
                         19961008
                    A1
                         19981021
                                       EP 1996-906024
    EP 872249
                                                        19960315
        R: CH, DE, FR, GB, LI, NL
                                      JP 1996-528274 19960315
    JP 3193057 B2 20010730
    US 5869306
                    A 19990209
                                       US 1997-913592 19970912
                    A 19950317
PRAI JP 1995-59261
    WO 1996-JP652
                    W
                         19960315
    A process for producing gene transfer prepns. by freeze-drying a mixt. of
AB
    a recombinant virus vector with at least one additive selected
    among arginine, glutamic acid (or sodium salt thereof), serine, glucose,
     inositol, lactose, mannitol, sorbitol, trehalose and xylose.
    The prepn. is to preserve the potency of the recombinant
    virus vectors. Prepn. of a compn. contq. recombinant MoMLV vector
    was shown.
L12 ANSWER 17 OF 41 CAPLUS COPYRIGHT 2003 ACS
    1995:640918 CAPLUS
AN
    123:29478
DN
TI
    Preservation of recombinant viruses by lyophilization
     in a stabilizing formulation
IN
    Herrmann, Steven M.; Prussak, Charles E.
    Viagene, Inc., USA
PΑ
SO
    PCT Int. Appl., 26 pp.
    CODEN: PIXXD2
DT
    Patent
    English
LΑ
FAN.CNT 1
                                      APPLICATION NO. DATE
    PATENT NO.
                   KIND DATE
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    WO 9510601 A1 19950420 WO 1994-US11414 19941007
PΤ
        W: AM, AU, BB, BG, BR, BY, CA, CN, CZ, FI, GE, HU, JP, KE, KG, KP,
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KR, KZ, LK, LT, LV, MD, MG, MN, MW, NO, NZ, PL, RO, RU, SD, SI,
            SK, TJ, TT, UA, UZ, VN
        RW: KE, MW, SD, SZ, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU,
            MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN,
            TD, TG
                           19950420
                                          CA 1994-2158935 19941007
     CA 2158935
                      AΑ
                           19950504
                                          AU 1994-79711
                                                           19941007
    AU 9479711
                      A1
                           19960828
                                          EP 1994-930662
                                                           19941007
    EP 728195
                      A1
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE
                          19970506
                                        JP 1994-511938
                                                         19941007
    JP 09504429
                      T2
PRAI US 1993-135938
                           19931012
    US 1993-153342
                           19931115
    WO 1994-US11414
                           19941007
AB
    Methods for preserving an infectious recombinant virus
     for subsequent reconstitution are provided. These methods are based upon
    methods of freeze-drying in the presence of stabilizers and
     cryoprotectants. The preferred method involves mixing the virus
    with an aq. soln. of a saccharide, a high mol. wt. structural additive, a
    buffering component and water to form an ag. suspension in which the
    virus is stabilized followed by cooling the aq. suspension to a
    temp. below the glass transition state temp. or below the eutectic point
     temp. of the formulation; and removing water from the cooled aq.
     suspension by sublimation to form a lyophilized virus having
     less than 10% water by wt. of the lyophilized virus. The
     virus is capable of infecting mammalian cells upon reconstitution.
     Optimization expts. are reported with conditions leading to storage with
     stable titers for up to 160 days reported.
    ANSWER 18 OF 41 CAPLUS COPYRIGHT 2003 ACS
L12
AN
     1993:456134 CAPLUS
DN
     119:56134
     Preservation of live bacteria with glycerol or
TI
    Clarke, Paul Douglas; Forrest, Bruce Darren
IN
PA
    Mastavac Ltd., UK
SO
     PCT Int. Appl., 17 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
    English
FAN.CNT 1
                                         APPLICATION NO. DATE
     PATENT NO.
                    KIND DATE
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                                          _____
    WO 9311220
                           19930610
                                         WO 1992-GB2243 19921203
PΙ
                     A1
        W: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP,
            KR, LK, LU, MG, MN, MW, NL, NO, PL, PT, RO, RU, SD, SE, UA, US
        RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE,
            BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, SN, TD, TG
                                         AU 1992-29546 19921203
    AU 9229546
                      A1
                           19930628
                           19911203
PRAI GB 1991-25695
    WO 1992-GB2243
                           19921203
    Live bacteria are mixed with trehalose (I) or glycerol
AB
     and dried to preserve them. An aq. suspension of Salmonella
     typhi contg. 2.3x1010 viable organism/mL and 20% I was coated on filter
    paper and air-dried. The no of viable cells after 7 days were 1x106 while
     there was no viable cell in the control contg. no I. Gastric resistant
    capsules contg. freeze- dried powder of S. typhi and I in an amt. to
    provide a viable count of 1-5x1010 cells were prepd.
L12 ANSWER 19 OF 41 CAPLUS COPYRIGHT 2003 ACS
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AN

DN

1954:78344 CAPLUS

48:78344

OREF 48:13823i,13824a The preservation of tomato spotted wilt virus by ΤI vacuum drying ΑU Finlay, K. W.; Parker, C. A. CS Univ. Western Australia, Nedlands J. Australian Inst. Agr. Sci. (1954), 20, 112-14 SO DT Journal LΑ Unavailable AB The virus survived vacuum drying when the infective plant sap was mixed with fresh egg albumin, but it was found necessary to include P2O5 in the ampuls to increase the longevity of the active virus. After storage for 21 months at room temp. the virus was still infective. L12 ANSWER 25 OF 41 CAPLUS COPYRIGHT 2003 ACS NA1940:39742 CAPLUS DN 34:39742 OREF 34:6021c-f Apparatus for the preservation of sera, bacterial cultures, viruses, etc. Flosdorf, Earl W. INTrustees of the University of Pennsylvania PA DT Patent LΑ Unavailable FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE -----US PIUS 2199815 19400507 Various structural and operative details are described, of app. for the ΆB treatment and preservation of biologically active substances by freezing the substance, dehydrating it from the frozen state under a high vacuum, and carrying out the operation in the final individual containers in which the resulting product is to be kept until used. U. S. 2,199,816 also describes numerous details of app. used and of operation for the treatment and preservation of biologically active substances by freezing the substance, dehydrating it from the frozen state under a high vacuum, and sealing the dried product under a high vacuum, the whole process being a continuous process and conducted under aseptic conditions in the final container, in which the material is to be stored, and distributed. U. S. 2,199,817 relates to the preservation of similar materials produced in a desiccated state in the container by freezing the substance, dehydrating it from the frozen state under a high vacuum, and sealing the dried product under a high vacuum, the container being provided with a rubber stopper which has a passage therethrough and an integral rubber tubular extension of said passage, the container being sealed by clamping off this rubber extension after the biologically active substance has been desiccated and while a vacuum is maintained within the container, by means or a metal clamp. Cf. C. A. 33, 8230.8. L12 ANSWER 18 OF 41 CAPLUS COPYRIGHT 2003 ACS AN 1993:456134 CAPLUS DN 119:56134 ΤI Preservation of live bacteria with glycerol or ΙN Clarke, Paul Douglas; Forrest, Bruce Darren PAMastavac Ltd., UK SO PCT Int. Appl., 17 pp. CODEN: PIXXD2

DT

Patent

English FAN.CNT 1 KIND DATE APPLICATION NO. DATE PATENT NO. ----------_____ WO 1992-GB2243 19921203 PΙ WO 9311220 A1 19930610 W: AT, AU, BB, BG, BR, CA, CH, CS, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, PL, PT, RO, RU, SD, SE, UA, US RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, SN, TD, TG 19930628 AU 1992-29546 AU 9229546 A1 PRAI GB 1991-25695 19911203 WO 1992-GB2243 19921203 Live bacteria are mixed with trehalose (I) or glycerol AB and dried to preserve them. An aq. suspension of Salmonella typhi contq. 2.3x1010 viable organism/mL and 20% I was coated on filter paper and air-dried. The no of viable cells after 7 days were 1x106 while there was no viable cell in the control contg. no I. Gastric resistant capsules contg. freeze- dried powder of S. typhi and I in an amt. to provide a viable count of 1-5x1010 cells were prepd. L12 ANSWER 33 OF 41 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V. 74162905 EMBASE AN DN 1974162905 TIA method for preservation of bacteria and bacteriophages by drying in vacuo. ΑU Iijima T.; Sakane T. CS Inst. Fermentat., Osaka, Japan Cryobiology, (1973) 10/5 (379-385). SO CODEN: CRYBAS DT Journal FS 004 Microbiology 047 Virology LΑ English AΒ An efficient and practical method was established to preserve bacterial strains and bacteriophages. The method is characterized by drying without freezing and by use of a cotton wool plug (nonabsorbent) to prevent contamination. Drying conditions were examined by measuring temperature, vacuum, and residual moisture of the samples. From measurement, it was found that the cotton wool plug acts as a buffer and a desiccant. Thus, the specimens reached optimal conditions during storage. Another advantage was that the temperature of the specimen during the drying procedure was 2-5.degree.C; therefore, the evaporation of the water is rapid and the time of completion is shorter than that during lyophilization. L12 ANSWER 36 OF 41 PROMT COPYRIGHT 2003 Gale Group AN 95:459400 PROMT ΤI Is sugar the secret of more stable drugs? SO Manufacturing Chemist, (Nov 1995) pp. 33. ISSN: 0262-4230. LΑ English WC 1405 *FULL TEXT IS AVAILABLE IN THE ALL FORMAT* AΒ Some scientists have predicted that, if ever a cure for cancer is to be discovered, it will be found deep in the rainforests. The plant world has already provided researchers with cures and treatments for many diseases. It invariably seems that the natural world can develop more bizarre and ingenious processes than the human mind is capable of dreaming up. For

example, the secret of eternal life is desired by many but beyond the

realms of science fiction B-movies it is an impossibility.

So is it conceivable that organisms exist with the ability to return to life after being dead for a hundred years, and that scientists are tapping into this phenomenon to develop products that could significantly alter healthcare around the world? Such organisms do exist, and they give the impression of being dried up and dead even when exposed to the most sensitive tests. But sprinkle them with water and, as if by magic, life returns.

This is a survival mechanism used by some plants and animals that live in the desert. In times of drought they dry out completely, but once the rains come, they are brought back to life. Known as cryptobionts, they can lose up to 99% of their water, remain dormant for long periods, and still return to full metabolic activity when rehydrated. There is one recorded case of a resurrection plant (Selaginella lepidophylla) which was restored after being kept in a museum for 120 years. Antonie van Leeuwenhoek first described the existence of cryptobionts in 1674, and over 300 years later a Cambridge scientist, Bruce Roser, founded the company Quadrant, to capitalise on this natural process. He wanted to prove that cryptobionts held the key to attaining the pharmaceutical goal of creating formulations stable at room temperature. So what is it about these organisms that means they can rise from the dead? The basis is a naturally-occurring disaccharide, trehalose. In order to survive the drought, a number of organisms such as the resurrection plant swap their water molecules for trehalose ones.

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=> d l14 1-34 ti,au, so

L14 ANSWER 1 OF 34 MEDLINE

DUPLICATE 1

- TI Re-suspension of T(1)44 vaccine cultures of Mycoplasma mycoides subsp mycoides SC in 1 molar MgSO(4) causes a drop in pH and a rapid reduction in titre.
- AU March John B; Waite Emma R; Litamoi Joseph K
- SO FEMS IMMUNOLOGY AND MEDICAL MICROBIOLOGY, (2002 Oct 11) 34 (2) 97-103. Journal code: 9315554. ISSN: 0928-8244.
- L14 ANSWER 2 OF 34 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 2
- TI Molecular epidemiology of contagious bovine pleuropneumonia in Tanzania based on amplified fragment length polymorphism and pulsed-field gel electrophoresis analysis
- AU Kusiluka, L. J. M.; Ojeniyi, B.; Friis, N. F.; Kokotovic, B.; Ahrens, P.
- SO Journal of Veterinary Medicine, Series B (2001), 48(4), 303-312 CODEN: JVMBE9: ISSN: 0931-1793
- L14 ANSWER 3 OF 34 MEDLINE

DUPLICATE 3

- TI Effect of HEPES buffer systems upon the pH, growth and survival of Mycoplasma mycoides subsp. mycoides small colony (MmmSC) vaccine cultures.
- AU Waite E R; March J B
- SO FEMS MICROBIOLOGY LETTERS, (2001 Jul 24) 201 (2) 291-4. Journal code: 7705721. ISSN: 0378-1097.
- L14 ANSWER 4 OF 34 MEDLINE

DUPLICATE 4

- TI Characterization of strains of Mycoplasma mycoides subsp.
 mycoides small colony type isolated from recent outbreaks of
 contagious bovine pleuropneumonia in Botswana
 and Tanzania: evidence for a new biotype.
- AU March J B; Clark J; Brodlie M
- SO JOURNAL OF CLINICAL MICROBIOLOGY, (2000 Apr) 38 (4) 1419-25. Journal code: 7505564. ISSN: 0095-1137.
- L14 ANSWER 5 OF 34 CAPLUS COPYRIGHT 2003 ACS
- TI Specific PCR identification of the T1 vaccine strains for contagious bovine pleuropneumonia
- AU Lorenzon, S.; David, A.; Nadew, M.; Wesonga, H.; Thiaucourt, F.
- SO Molecular and Cellular Probes (2000), 14(4), 205-210 CODEN: MCPRE6; ISSN: 0890-8508
- L14 ANSWER 6 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE
- TI Inhibitory and mycoplasmacidal concentrations of some antibiotics on strains of Mycoplasma mycoides subsp. mycoides SC: The causative agent of contagious bovine pleuropneumonia.
- AU Egwu, G. O.; Aliyu, M. M.
- SO Acta Veterinaria (Belgrade), (1998) Vol. 48, No. 5-6, pp. 309-315. ISSN: 0567-8315.
- L14 ANSWER 7 OF 34 MEDLINE

DUPLICATE 6

- TI Contagious bovine pleuropneumonia. The costs of control in Central/southern Africa.
- AU Windsor R S; Wood A
- SO ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1998 Jun 29) 849 299-306. Journal code: 7506858. ISSN: 0077-8923.
- L14 ANSWER 8 OF 34 MEDLINE

DUPLICATE 7

TI Vaccination against contagious bovine pleuropneumonia and the use of molecular tools in epidemiology.

- AU Thiaucourt F; Lorenzon S; David A; Tulasne J J; Domenech J
- SO ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1998 Jun 29) 849 146-51.

 Ref: 15

 Journal code: 7506858. ISSN: 0077-8923.
- L14 ANSWER 9 OF 34 MEDLINE

DUPLICATE 8

- TI ISCOM vaccine against contagious bovine pleuropneumonia (CBPP).

 1. Biochemical and immunological characterization.
- AU Abusugra I; Wolf G; Bolske G; Thiaucourt F; Morein B
- SO VETERINARY IMMUNOLOGY AND IMMUNOPATHOLOGY, (1997 Oct 6) 59 (1-2) 31-48. Journal code: 8002006. ISSN: 0165-2427.
- L14 ANSWER 10 OF 34 CAPLUS COPYRIGHT 2003 ACS
- Polymerase chain reaction (PCR) assay for the detection and differentiation of the virulent strains of Mycoplasma mycoides subspecies mycoides "small colony" type from the vaccine t1 strain in purified DNA and in field samples from Africa (cytadhesin, P1, contagious bovine pleuropneumonia)
- AU Kalif, Abdullahi Sheikh Hassan
- SO (1996) 97 pp. Avail.: UMI, Order No. DA9720360 From: Diss. Abstr. Int., B 1997, 58(2), 465
- L14 ANSWER 11 OF 34 MEDLINE
- TI [Strategies for prevention and eradication of contagious bovine pleuropneumonia with or without vaccination].

 Strategies de prophylaxie et d'eradication de la peripneumonie contagieuse bovine avec ou sans vaccination.
- AU Provost A
- SO REVUE SCIENTIFIQUE ET TECHNIQUE, (1996 Dec) 15 (4) 1355-71. Ref: 27 Journal code: 8712301. ISSN: 0253-1933.
- L14 ANSWER 12 OF 34 MEDLINE

DUPLICATE 9

- Monoclonal antibodies to surface-exposes proteins of Mycoplasma mycoides subsp. mycoides (small-colony strain), which causes contagious bovine pleuropneumonia.
- AU Kiarie M N; Rurangirwa F R; Perryman L E; Jasmer D P; McGuire T C
- SO CLINICAL AND DIAGNOSTIC LABORATORY IMMUNOLOGY, (1996 Nov) 3 (6) 746-52. Journal code: 9421292. ISSN: 1071-412X.
- L14 ANSWER 13 OF 34 MEDLINE

DUPLICATE 10

- TI Insertion element IS1296 in Mycoplasma mycoides subsp. mycoides small colony identifies a European clonal line distinct from African and Australian strains.
- AU Cheng X; Nicolet J; Poumarat F; Regalla J; Thiaucourt F; Frey J
- SO MICROBIOLOGY, (1995 Dec) 141 (Pt 12) 3221-8. Journal code: 9430468. ISSN: 1350-0872.
- L14 ANSWER 14 OF 34 MEDLINE
- TI Contagious bovine pleuropneumonia **vaccines**: the need for improvements.
- AU Rweyemamu M M; Litamoi J; Palya V; Sylla D
- SO REVUE SCIENTIFIQUE ET TECHNIQUE, (1995 Sep) 14 (3) 593-601. Ref: 14 Journal code: 8712301. ISSN: 0253-1933.
- L14 ANSWER 15 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI THE EFFECT OF RECONSTITUTING DILUENTS ON THE VIABILITY OF T-1 STRAIN MYCOPLASMA VACCINE.
- AU GARBA S A; NGBEDE J
- SO DISCOVERY INNOVATION, (1991) 3 (1), 63-65. CODEN: DIINE4. ISSN: 1015-079X.

- L14 ANSWER 16 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI SEROSURVEY OF SAHELIAN CATTLE FOR EVIDENCE OF EPIZOOTIC DISEASE.
- AU MARINER J C; SAMA S; MAMINI C; BAARE K; STEM C; YEDLOUTSCHNIG R J; MEBUS C A; SOLLOD A E
- SO PREV VET MED, (1989) 7 (3), 163-172. CODEN: PVMEEG.
- L14 ANSWER 17 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI PROSPECTS OF MYCOPLASMA VACCINES WITH SPECIAL EMPHASIS ON OIL-BASED INACTIVATED VACCINES.
- AU GARBA S A
- SO SIXTH INTERNATIONAL CONGRESS OF THE INTERNATIONAL ORGANIZATION FOR MYCOPLASMOLOGY, BIRMINGHAM, ALABAMA, USA, AUGUST 26-31, 1986. ISR J MED SCI. (1987) 23 (5), 535.

 CODEN: IJMDAI. ISSN: 0021-2180.
- L14 ANSWER 18 OF 34 MEDLINE
- TI A reassessment of the dual **vaccine** against rinderpest and contagious bovine pleuropneumonia.
- AU Jeggo M H; Wardley R C; Corteyn A H
- SO VETERINARY RECORD, (1987 Feb 7) 120 (6) 131-5. Journal code: 0031164. ISSN: 0042-4900.
- L14 ANSWER 19 OF 34 MEDLINE DUPLICATE 11
- TI Observations on experimental inactivated **vaccines** for contagious bovine pleuropneumonia.
- AU Gray M A; Simam P; Smith G R
- SO JOURNAL OF HYGIENE, (1986 Oct) 97 (2) 305-15. Journal code: 0375374. ISSN: 0022-1724.
- L14 ANSWER 20 OF 34 SCISEARCH COPYRIGHT 2003 ISI (R)
- TI MYCOPLASMA VACCINES THE POTENCY OF WET T1 BROTH CONTAGIOUS BOVINE PLEUROPNEUMONIA (CBPP)
 VACCINE IN CATTLE AFTER STORAGE AT 4-DEGREES-C
- AU GARBA S A (Reprint); IMOGIE U A
- SO YALE JOURNAL OF BIOLOGY AND MEDICINE, (1984) Vol. 57, No. 6, pp. 904.
- L14 ANSWER 21 OF 34 MEDLINE
- TI The ability of Mycoplasma mycoides subspecies mycoides and closely related strains from goats and sheep to immunize mice against subspecies capri.
- AU Smith G R; Oliphant J C
- SO JOURNAL OF HYGIENE, (1981 Oct) 87 (2) 321-9. Journal code: 0375374. ISSN: 0022-1724.
- L14 ANSWER 22 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI IMMUNOGENIC STABILITY OF HEAT KILLED MYCOPLASMA-MYCOIDES-SSP-MYCOIDES AT 4 CELSIUS.
- AU HOOKER J M; SMITH G R; MILLIGAN R A
- SO BR VET J, (1980) 136 (6), 614-616. CODEN: BVJOA9. ISSN: 0007-1935.
- L14 ANSWER 23 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 12
- TI IMMUNE RESPONSE OF MICE RABBITS AND CATTLE TO INACTIVATED MYCOPLASMA-MYCOIDES-SSP-MYCOIDES VACCINES CONTAINING ADJUVANTS.
- AU HOOKER J M; SMITH G R; MILLIGAN R A
- SO J COMP PATHOL, (1980) 90 (3), 363-372. CODEN: JCVPAR. ISSN: 0021-9975.
- L14 ANSWER 24 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI PATHOGENICITY OF A CAPRINE STRAIN OF MYCOPLASMA-MYCOIDES-SSP-MYCOIDES FOR

CATTLE.

AU OJO M O; KASALI O B; OZOYA S E

SO J COMP PATHOL, (1980) 90 (2), 209-216. CODEN: JCVPAR. ISSN: 0021-9975.

- L14 ANSWER 25 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI SOME EVIDENCE OF AN AGE SUSCEPTIBILITY TO CONTAGIOUS BOVINE PLEURO PNEUMONIA.
- AU MASIGA W N; WINDSOR R S
- SO RES VET SCI, (1978) 24 (3), 328-333. CODEN: RVTSA9. ISSN: 0034-5288.
- L14 ANSWER 26 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- TI CONTAGIOUS BOVINE PLEURO PNEUMONIA COMPARATIVE EFFICACY TRIAL OF THE FREEZE DRIED FRENCH T-1 VACCINE AND THE T-1 BROTH CULTURE VACCINE MUGUGA.
- AU MASIGA W N; RURANGIRWA F R; ROBERTS D H; KAKOMA I
- SO BULL ANIM HEALTH PROD AFR, (1978 (RECD 1979)) 26 (3), 216-223. CODEN: BAHADH. ISSN: 0378-9721.
- L14 ANSWER 27 OF 34 MEDLINE
- TI An investigation into the viability of broth cultures of the T1 strain of Mycoplasma mycoides sub-species mycoides.
- AU Windsor R S
- SO RESEARCH IN VETERINARY SCIENCE, (1978 Jan) 24 (1) 109-12. Journal code: 0401300. ISSN: 0034-5288.
- L14 ANSWER 28 OF 34 MEDLINE DUPLICATE 13
- TI Virulence of established **vaccine** strains and artificially passaged field strain of Mycoplasma mycoides subsp mycoides.
- AU Dyson D A; Smith G R
- SO RESEARCH IN VETERINARY SCIENCE, (1976 Mar) 20 (2) 185-90. Journal code: 0401300. ISSN: 0034-5288.
- L14 ANSWER 29 OF 34 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI The effect of post vaccinal treatment with the antibiotic tylosin on the immunity produced by the Tl strain of Mycoplasma mycoides subspecies Mycoides.
- AU Windsor R.S.; Masiga W.N.
- SO Journal of Comparative Pathology, (1976) 86/2 (173-181). CODEN: JCVPAR
- L14 ANSWER 30 OF 34 MEDLINE
- TI Contagious bovine pleuropneumonia-protection following natural infection and vaccination.
- AU Gourlay R N
- SO DEVELOPMENTS IN BIOLOGICAL STANDARDIZATION, (1975) 28 586-9. Journal code: 0427140. ISSN: 0301-5149.
- L14 ANSWER 31 OF 34 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI Attempts to differentiate Mycoplasma mycoides var. mycoides immune cattle from susceptible cattle.
- AU Roberts D.H.; Windsor R.S.
- SO Research in Veterinary Science, (1974) 17/3 (403-405). CODEN: RVTSA
- L14 ANSWER 32 OF 34 MEDLINE DUPLICATE 14
- TI Factors affecting the viability of Mycoplasma mycoides in bottled contagious bovine pleuropneumonia vaccine
- AU Lloyd L C; Pearson C W; Etheridge J R

- SO JOURNAL OF APPLIED BACTERIOLOGY, (1974 Sep) 37 (3) 297-307. Journal code: 7503050. ISSN: 0021-8847.
- L14 ANSWER 33 OF 34 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI Immunoelectrophoretic analysis of Mycoplasma mycoides var. mycoides.
- AU Stone S.S.; Razin S.
- SO Infection and Immunity, (1973) 7/6 (922-930). CODEN: INFIBR
- L14 ANSWER 34 OF 34 MEDLINE
- TI The immunizing dose of T1 strain Mycoplasma mycoides against contagious bovine pleuropneumonia.
- AU Gilbert F R; Windsor R S
- SO TROPICAL ANIMAL HEALTH AND PRODUCTION, (1971) 3 (2) 71-6. Journal code: 1277355. ISSN: 0049-4747.

ANSWER 8 OF 34 DUPLICATE 7 MEDLINE L14MEDLINE AN 1998333056 98333056 PubMed ID: 9668459 DN Vaccination against contagious bovine pleuropneumonia and the use of TImolecular tools in epidemiology. Thiaucourt F; Lorenzon S; David A; Tulasne J J; Domenech J ΑU CIRAD-EMVT, Montpellier, France. CS ANNALS OF THE NEW YORK ACADEMY OF SCIENCES, (1998 Jun 29) 849 146-51. SO Journal code: 7506858. ISSN: 0077-8923. CY United States Journal; Article; (JOURNAL ARTICLE) DT General Review; (REVIEW) (REVIEW, TUTORIAL) LΑ English FS Priority Journals EΜ 199808 ΕĎ Entered STN: 19980820 Last Updated on STN: 19980820 Entered Medline: 19980810 Contagious bovine pleuropneumonia is a serious threat to cattle not only in Africa but also in southern Europe and possibly Asia. It is now present in countries that had been free of the disease for many years, giving rise to doubts about the efficiency of the control strategies. In Africa CBPP is controlled mainly by a vaccination policy that uses variant strains of Mycoplasma mycoides subsp mycoides biotype SC, called T1/44 or Tlsr. Until recently, it was not possible to differentiate the various strains within the biotype and consequently to identify the vaccine strains. Restriction analysis of mycoplasma DNA with HindIII and Pst1 has been applied to 24 strains of African origin and one European strain. Each enzyme gave rise to different restriction profiles and the combination of the results permitted subdivision of these strains into 9 groups. Interestingly, some profiles of pathogenic strains seem to be restricted to certain geographical areas. The profile of the poorly immunogenic vaccinal strain KH3J is also very peculiar, and it is easily distinguished from that of the other vaccine strains originating from T1. This technique is simple once the strains are isolated. Efforts are now under way to use molecular tools based on PCR products to alleviate the difficulty of isolation. DUPLICATE 8 ANSWER 9 OF 34 MEDLINE L14AN1998100478 MEDLINE DN 98100478 PubMed ID: 9437824 ISCOM vaccine against contagious bovine pleuropneumonia (CBPP). TI 1. Biochemical and immunological characterization. ΑU Abusugra I; Wolf G; Bolske G; Thiaucourt F; Morein B

Swedish University of Agricultural Sciences, Faculty of Veterinary

Medicine, Department of Microbiology, Uppsala, Sweden..

SO VETERINARY IMMUNOLOGY AND IMMUNOPATHOLOGY, (1997 Oct 6) 59 (1-2) 31-48. Journal code: 8002006. ISSN: 0165-2427.

DT Journal; Article; (JOURNAL ARTICLE)

izzeldin.abusugra@bmc.uu.se

LA English

CS

CY

FS Priority Journals

Netherlands

EM 199802

ED Entered STN: 19980224

Last Updated on STN: 19980224 Entered Medline: 19980212 AB A better vaccine than the existing ones against contagious bovine pleuropneumonia (CBPP)

caused by Mycoplasma mycoides subsp. mycoides small colony type (MmmSC) would improve the chances for eradication of CBPP. In such an effort, immunostimulating complexes (ISCOMS) have been prepared from the whole detergent-solubilized cells of MmmSC and characterized biochemically and immunologically. The most efficient detergent for solubilization of the mycoplasma was MEGA-10 which yielded a high recovery of proteins in the ISCOMS. The ISCOMS showed the typical cage-like structure by EM and sedimented as 19S by sucrose gradient centrifugation. The protein pattern of the ISCOMS, analyzed in SDS-PAGE, revealed a great number of bands distributed along the gel as high and low molecular weight polypeptides. The Western blot developed with a serum from a CBPP infected animal detected a reduced number of polypeptides. In samples from whole mycoplasma cells and in ISCOMS, lectin blots revealed more than 20 carbohydrate structures. The ISCOMS induced a strong primary antibody response in mice measured by ELISA and the boost resulted in a 6-fold increase of the serum antibody response. The IqG response was distributed into various IgG subclasses with high IgG1, IgG2a and IgG2b titres while the IgG3 response was low. In cattle the ISCOM vaccine induced strong primary and long lasting secondary antibody responses of similar magnitudes as those of naturally infected animals as recorded by ELISA which persisted more than a year. IgG response was equally distributed in IqG1 and IqG2 subclasses. Also a cell-mediated immune response measured by proliferation assay was induced by low dose of ISCOMS. In the growth inhibition test, sera from vaccinated cattle readily inhibited colony growth already after the first immunization.

L14 ANSWER 12 OF 34 MEDLINE

DUPLICATE 9

AN 97071924 MEDLINE

DN 97071924 PubMed ID: 8914769

DN 9/0/1924 PubMed 1D: 8914/69

- TI Monoclonal antibodies to surface-exposes proteins of Mycoplasma mycoides subsp. mycoides (small-colony strain), which causes contagious bovine pleuropneumonia.
- AU Kiarie M N; Rurangirwa F R; Perryman L E; Jasmer D P; McGuire T C
- CS. Department of Veterinary Microbiology and Pathology, Washington State University, Pullman 99164-7040, USA.
- SO CLINICAL AND DIAGNOSTIC LABORATORY IMMUNOLOGY, (1996 Nov) 3 (6) 746-52. Journal code: 9421292. ISSN: 1071-412X.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 199702
- ED Entered STN: 19970305

Last Updated on STN: 19970305

Entered Medline: 19970218

Outbreaks of bovine pleuropneumonia caused by small-colony strains of Mycoplasma mycoides subsp. mycoides occur in Africa, and vaccination is used for control. Since protein subunits are needed to improve multivalent vaccines, monoclonal antibodies (MAbs) were made to facilitate protein identification and isolation. Eleven immunoglobulin M MAbs derived from mouse spleen donors immunized with disrupted whole organisms bound periodate-sensitive epitopes on externally exposed polysaccharide. Seven of these MAbs caused in vitro growth inhibition of M. mycoides subsp. mycoides; however, reaction with carbohydrate epitopes prevented their use in identifying proteins. Ten additional MAbs from mouse spleen donors immunized with Triton X-114-phase integral membrane proteins reacted with periodate-insensitive, proteinase K-sensitive epitopes. These MAbs were classified into three groups based on immunoblots of Triton X-114-phase proteins. One group reacted with 96-, 16-, and 15-kDa proteins. Another

group reacted with 26-, 21-, and 16-kDa proteins, while a third group reacted only with 26- and 21-kDa proteins. One MAb from each group reacted with trypsinsensitive epitopes on live organisms, yet none caused in vitro growth inhibition. Representative MAbs reacted with all small-colony strains in immunoblots and did not react with large colony strains. However, these MAbs were not specific for small-colony strains, as proteins from two other M. mycoides cluster organisms were identified. Nevertheless, MAbs to surface-exposed epitopes on integral membrane proteins will be useful for isolation of these proteins for immunization, since one or more might induce growth-inhibiting antibodies or other protective responses.

L14 ANSWER 14 OF 34 MEDLINE

ΑN 96014310 MEDLINE

PubMed ID: 8593393 DN

- ΤI Contagious bovine pleuropneumonia vaccines: the need for improvements.
- ΑU Rweyemamu M M; Litamoi J; Palya V; Sylla D
- Food and Agriculture Organisation of the United Nations, Animal Health CS Service, Rome, Italy.
- REVUE SCIENTIFIQUE ET TECHNIQUE, (1995 Sep) 14 (3) 593-601. Ref: 14 SO Journal code: 8712301. ISSN: 0253-1933.
- CY France
- DT Journal; Article; (JOURNAL ARTICLE) General Review; (REVIEW) (REVIEW, TUTORIAL)
- LΑ English
- FS Priority Journals
- EM199604
- ED Entered STN: 19960422 Last Updated on STN: 19960422 Entered Medline: 19960411
- Contagious bovine pleuropneumonia (CBPP) AB

vaccines are routinely used only in Africa. The vaccines are usually produced from one of two strains (T1/44 and KH3J), each of which has a streptomycin-resistant variant. The necessity for a 'master seed strain' is evident. At least one manufacturer in Africa produces a broth culture vaccine, while others produce a freeze-dried product. A standardised manufacturing protocol needs to be developed, together with in-process and final product quality control procedures. Some CBPP vaccine manufacturing procedures do not allow sufficient leeway for the execution of typical quality control practices. For example, it is difficult to perform batch testing on broth culture vaccine, as the vaccine is produced in its final container. Quality control test results from the Pan African Veterinary Vaccine Centre (PANVAC) are analysed in terms of causes of batch failure and indicators for process development. Taking potency as an example, most vaccine batches tested by PANVAC pass only at the limit of the OIE minimum requirement of 10(7) colony-forming units per dose. To improve the titre of the vaccine, it will be necessary to modify the manufacturing process, either by increasing mycoplasma yield during the culture phase or by minimising losses during downstream processes, especially freeze-drying. Data on inactivated vaccines are scarce. Duration of the immunity achieved with live CBPP vaccines is relatively short, in comparison with other live vaccines. Data may be required on the molecular basis of virulence and immunogenicity, as well as on the molecular immunology of CBPP, to enable the development of improved vaccines.

L14 ANSWER 15 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

AN 1992:6347 BIOSIS DN BA93:6347

TI THE EFFECT OF RECONSTITUTING DILUENTS ON THE VIABILITY OF T-1 STRAIN MYCOPLASMA VACCINE.

AU GARBA S A; NGBEDE J

- CS DEP. BIOLOGICAL SCIENCES, FEDERAL UNIVERSITY TECHNOLOGY, P.M.B. 65 MINNA, NIGERIA.
- SO DISCOVERY INNOVATION, (1991) 3 (1), 63-65. CODEN: DIINE4. ISSN: 1015-079X.
- FS BA; OLD
- LA English
- AB Six diluents were used to reconstitute 12 batches of freeze-dried T1 strain contagious bovine pleuropneumonia
 vaccine. From the findings, normal saline is the most suitable diluent as it stored well at 4.degree. C for 10 hours with a drop of viable mycoplasma from 2.7 .times. 108 cfu/ml to 2.2 .times. 108 cfu/ml. The order of performance of these diluents was: normal saline > distilled water > well water > spring water > dam water > tap water. The vaccine stored at 42.degree. C deteriorated rapidly within 10 hours of reconstitution with a performance pattern similar to those stored at 4.degree. C.
- L14 ANSWER 17 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AN 1988:216430 BIOSIS
- DN BR34:109440
- TI PROSPECTS OF MYCOPLASMA VACCINES WITH SPECIAL EMPHASIS ON OIL-BASED INACTIVATED VACCINES.
- AU GARBA S A
- CS NATL. VET. RES. INST., VOM, NIGERIA.
- SO SIXTH INTERNATIONAL CONGRESS OF THE INTERNATIONAL ORGANIZATION FOR MYCOPLASMOLOGY, BIRMINGHAM, ALABAMA, USA, AUGUST 26-31, 1986. ISR J MED SCI. (1987) 23 (5), 535.

 CODEN: IJMDAI. ISSN: 0021-2180.
- DT Conference
- FS BR; OLD
- LA English
- L14 ANSWER 19 OF 34 MEDLINE

DUPLICATE 11

- AN 87058903 MEDLINE
- DN 87058903 PubMed ID: 3782785
- TI Observations on experimental inactivated vaccines for contagious bovine pleuropneumonia.
- AU Gray M A; Simam P; Smith G R
- SO JOURNAL OF HYGIENE, (1986 Oct) 97 (2) 305-15. Journal code: 0375374. ISSN: 0022-1724.
- CY ENGLAND: United Kingdom
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 198612
- ED Entered STN: 19900302

Last Updated on STN: 19900302

Entered Medline: 19861224

AB In two trials the efficacy of inactivated vaccines against contagious bovine pleuropneumonia was tested by exposing vaccinated cattle to droplet infection provided by close contact with experimentally infected 'donors'. Complete protection was given by an extreme form of vaccination in which a heavy suspension of killed Mycoplasma mycoides subsp. mycoides emulsified with Freund's complete adjuvant was given in two large doses. 'Mouse-protective antibody' (MPA) was also produced, i.e. serum transferred to mice 2-4 h before intraperitoneal challenge prevented the development of

mycoplasmaemia. However, the study did not answer the question 'Is MPA protective for cattle?'. No protection was given by a milder form of vaccination in which a lighter suspension of killed mycoplasmas emulsified with Freund's incomplete adjuvant was given in a comparatively small dose on a single occasion.

L14 ANSWER 21 OF 34 MEDLINE

AN 82031798 MEDLINE

DN 82031798 PubMed ID: 7026674

TI The ability of Mycoplasma mycoides subspecies mycoides and closely related strains from goats and sheep to immunize mice against subspecies capri.

AU Smith G R; Oliphant J C

SO JOURNAL OF HYGIENE, (1981 Oct) 87 (2) 321-9. Journal code: 0375374. ISSN: 0022-1724.

CY ENGLAND: United Kingdom

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 198112

ED Entered STN: 19900316 Last Updated on STN: 19900316

Entered Medline: 19811215

AB Small colony (SC) strains of Mycoplasma mycoides subsp. mycoides from contagious bovine pleuropneumonia (CBPP) and from goats were compared with large colony (LC) strains of so-called M. mycoides subsp. mycoides from goats and sheep by means of a cross-protection test in which mice were challenged with M. mycoides subsp. capri. Of 13 LC strains, all gave partial cross-protection, and 11 were shown to be more closely related than four SC strains to subspecies capri. In a further experiment, six SC strains--three from CBPP and three from goats--all gave weak partial cross-protection against subspecies capri.

- L14 ANSWER 23 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.DUPLICATE 12
- AN 1981:132871 BIOSIS
- DN BA71:2863
- TI IMMUNE RESPONSE OF MICE RABBITS AND CATTLE TO INACTIVATED
 MYCOPLASMA-MYCOIDES-SSP-MYCOIDES VACCINES CONTAINING ADJUVANTS.
- AU HOOKER J M; SMITH G R; MILLIGAN R A
- CS NUFFIELD LAB. COMP. MED., INST. ZOOL., ZOOL. SOC. LOND., REGENT'S PARK, LONDON NW1 4RY, ENGL., UK.
- SO J COMP PATHOL, (1980) 90 (3), 363-372. CODEN: JCVPAR. ISSN: 0021-9975.
- FS BA; OLD
- LA English
- AΒ Mice, rabbits and cattle were inoculated s.c. with various dilutions of a suspension of heat-killed M. mycoides ssp. mycoides emulsified with equal volumes of Freund's incomplete adjuvant (FIA) or an Arachis oil adjuvant (AOA). The vaccine dose volumes for mice, rabbits and cattle were 0.2, 0.5 and 5.0 ml, respectively. The immune response of mice was measured by direct challenge; the responses of rabbits and cattle were measured by a passive mouse protection test. The response in rabbits and cattle was much greater than that in mice. The efficacy of FIA was much greater than that of AOA. A Mycoplasma suspension with an optical opacity equivalent to that of Brown's tube 2 emulsified with FIA produced a response in all 3 animal species within 6 wk. In rabbits given a 2nd dose of FIA vaccine or AOA vaccine 6-8 wk after the 1st, mouse-protective antibody (MPA) titers did not increase significantly. Demonstrable levels of MPA persisted in cattle and rabbits for several months after inoculation with FIA vaccine, and in

some animals for 10-12 mo. The titers in rabbits were sometimes very high. Further studies on the development of an inactivated **vaccine** for **contagious bovine pleuropneumonia** should be done.

- L14 ANSWER 26 OF 34 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AN 1979:238792 BIOSIS
- DN BA68:41296
- TI CONTAGIOUS BOVINE PLEURO PNEUMONIA COMPARATIVE EFFICACY TRIAL OF THE FREEZE DRIED FRENCH T-1 VACCINE AND THE T-1 BROTH CULTURE VACCINE MUGUGA.
- AU MASIGA W N; RURANGIRWA F R; ROBERTS D H; KAKOMA I
- CS VET. RES. ORGAN., P.O. BOX 32, MUGUGA, KIKUYU, KENYA.
- SO BULL ANIM HEALTH PROD AFR, (1978 (RECD 1979)) 26 (3), 216-223. CODEN: BAHADH. ISSN: 0378-9721.
- FS BA; OLD
- LA English
- Except for 2 animals, all the animals in the group vaccinated with the AΒ Tiper T1 freeze-dried vaccine and challenged 6 mo. after vaccination had a serological response during the challenge period. Except for 3 animals, all the animals in the group vaccinated with Tl broth culture vaccine and challenged 6 mo. later developed CF [complement-fixing] antibodies [Ab]. Four animals vaccinated with the T1 broth culture vaccine had CF Ab titers during the trial performed 15 mo. after primary vaccination. Of 15 animals vaccinated with Tiper T1 freeze-dried vaccine and challenged 15 mo. after primary vaccination, 12 had CF Ab titers during the trial. Of 25 control animals in the 15-mo. trial, 19 had a serological response. No CBPP [contagious bovine pleuropneumonial lesions were present in the animals vaccinated with Tiper T1 freeze-dried ${\bf vaccine}$ or T1 broth culture ${\bf vaccine}$ and challenged 6 mo. after primary vaccination. Of the control animals in the 6-mo. trial, 12 died of other causes; the remaining 10 animals were killed 6 wk after the onset of the CF Ab. Mycoplasma mycoides was recovered from 28 of 35 control animals. Four animals which were vaccinated with T1 broth culture vaccine and challenged 15 mo. later had CBPP lesions at post-mortem examination. Of 15 of the animals vaccinated with Tiper T1 freeze-dried vaccine, 7 had CBPP lesions at post-mortem examination. Of 25 control animals in the trial performed 15 mo. post-primary vaccination, 11 died of CBPP and 4 animals were killed in extremis.